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Substitute for form 1449A/PTO SUPPLEMENTAL INFORMATION DISCLOSURE STATEMENT BY APPLICANT <i>(use as many sheets as necessary)</i>		Complete if Known <table border="1" style="width: 100%; border-collapse: collapse;"> <tr> <td style="width: 50%;">Application Number</td> <td>09/834,596</td> </tr> <tr> <td>Filing Date</td> <td>April 13, 2001</td> </tr> <tr> <td>First Named Inventor</td> <td>Watanabe <i>et al.</i></td> </tr> <tr> <td>Group Art Unit</td> <td>1623</td> </tr> <tr> <td>Examiner Name</td> <td>Howard V. Owens, Jr.</td> </tr> <tr> <td>Attorney Docket Number</td> <td>08841.105037 PHAR 2020</td> </tr> </table>		Application Number	09/834,596	Filing Date	April 13, 2001	First Named Inventor	Watanabe <i>et al.</i>	Group Art Unit	1623	Examiner Name	Howard V. Owens, Jr.	Attorney Docket Number	08841.105037 PHAR 2020
Application Number	09/834,596														
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Attorney Docket Number	08841.105037 PHAR 2020														
Sheet	1	of	7												

3372258_3

U.S. PATENT DOCUMENTS						
Examiner Initials *	Cite No. ¹	U.S. Patent Document		Name of Patentee or Applicant of Cited Document	Date of Publication of Cited Document MM-DD-YYYY	Pages, Columns, Lines, Where Relevant Passages/Relevant Figures Appear
		Number	Kind Code (if known)			
<i>Ho</i>	AA	3,480,613		Walton <i>et al.</i>	11-25-1969	
	AB	5,977,061		Holy <i>et al.</i>	11-02-1999	
	AC	6,340,690	B1	Bachand <i>et al.</i> (Idenix Pharm.)	01-22-2002	
	AD	6,395,716	B1	Gosselin <i>et al.</i> (Idenix Pharm.)	05-28-2002	
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	AF	6,573,248	B2	Ramasamy <i>et al.</i>	06-03-2003	
	AG	2002/0055483	A1	Watanabe <i>et al.</i>	05-09-2002	
	AH	2002/0147160	A1	Bhat <i>et al.</i>	10-10-2002	
	AI	2003/0008841	A1	Devos <i>et al.</i>	01-09-2003	
	AJ	2003/0028013	A1	Wang <i>et al.</i>	02-06-2003	
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<i>Ho</i>	AM	2003/0087873	A1	Stuyver <i>et al.</i>	05-08-2003	

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		Office ³	Number	Kind Code ² (if known)				
	AN	FR	1,521,076	✓	Merck & Co. Inc.	04-12-1968		
	AO	FR	1,581,628	✓	Merck & Co. Inc.	09-19-1969		
	AP	FR	2,662,165	✓	A1 Univ Pierre et Marie Curie, Paris	11-22-1991		
Ho	AQ	GB	1,163,103	A ✓	Merck & Co. Inc.	09-04-1969		
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	AS	JP	63-215694	✓	A Yamasa Shoyu Co. Ltd.	09-08-1988		
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Ho	AU	WO	98/16184	✓	A2 ICN Pharmaceuticals Inc.	04-23-1998		
Ho	AV	WO	99/43691	✓	A1 Emory U./Georgia Res. Found.	09-02-1999		

Examiner Signature <i>Howard V. Owens, Jr.</i>	Date Considered <i>6/30/04</i>
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Sheet 2 of 7

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		Office ³	Number	Kind Code ² (if known)				
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	BB	WO	01/16671	A1	Novirio (Idenix Pharmaceuticals)	03-08-2001		
	BC	WO	01/32153	A2	Biochem Pharma, Inc.	05-10-2001		
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	BK	WO	02/18404	A2	F. Hoffmann-La Roche AG	03-07-2002		
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	BM	WO	02/48165	A2	Pharmasset Ltd.	06-20-2002		
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X/10	BZ	WO	03/062257	A1	Ribapharm Inc.	07-31-2003		

Examiner Signature

Howard V. Owens, Jr.

Date Considered

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Sheet	3	of	7
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		Office ³	Number	Kind Code ² (if known)				
HO	CA	WO	03/063771	A2	Pharmasset Ltd.	08-07-2003		
HO	CB	WO	03/068162	A2	Pharmasset Ltd.	08-21-2003		
HO	CC	WO	03/072757	A2	Biota Inc.	09-04-2003		
HO	CD	WO	03/093290	A2	Genelabs Technologies Inc.	11-13-2003		

OTHER PRIOR ART - NON PATENT LITERATURE DOCUMENTS

Examiner Initials *	Cite No. †	Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine, journal, serial, symposium, catalog, etc.), date, page(s), volume-issue number(s), publisher, city and/or country where published.	T*
	CE	ALTMANN, K.H., <i>et al.</i> , "The Synthesis of 1'-methylcarboecyclic thymidine and its effect on nucleic acid duplex stability", <i>Synlett</i> , Thieme Verlag, Stuttgart, Gc, October 1994, 10, 853-855	
	CF	BAGINSKY, S.G. <i>et al.</i> , "Mechanism of action of a pestivirus antiviral compound," <i>Proc. Nat. Acad. Sci. (USA)</i> 2000, 97(14), 7981-7986.	
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Harold C. Co.

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Sheet 4 of 7

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OTHER PRIOR ART - NON PATENT LITERATURE DOCUMENTS

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Sheet **5** of **7**

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	EA	KAWANA, M., et al., "The deoxygenation of tosylated adenosine derivatives with Grignard reagents," <i>Nucleic Acids Symp. Ser.</i>, 1986, 17, 37-40.	
	EB	LAVARE, S., et al., "3'-deoxy-3'-trifluoromethyl nucleosides: synthesis and antiviral evaluation," <i>Nucleosides & Nucleotides</i>, 1998, 17, 2267-2280.	
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	EI	MIKHAILOV, S.N., et al., "Synthesis and properties of 3'-C-methylnucleosides and their phosphoric esters," <i>Carbohydrate Research</i>, 1983, 124, 75-96.	
	EJ	MIKHAILOV, S.N., et al., "Hydrolysis of 2'- and 3'-C-methyluridine 2',3'-cyclic monophosphates and interconversion and dephosphorylation of the resulting 2'- and 3'-monophosphates: comparison with the reactions of uridine monophosphates," <i>J. Org. Chem.</i>, 1992, 57, 4122-4126.	
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	EL	NUTT, R.F., et al., "Branched-chain sugar nucleosides. III. 3'-C-methyladenine," <i>J. Org. Chem.</i> 1968, 33, 1789-1795.	

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First Named Inventor	Watanabe <i>et al.</i>
Group Art Unit	1623
Examiner Name	Howard V. Owens, Jr.
Attorney Docket Number	08841.105037 PHAR 2020

3372258 3

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Examiner Initials *	Cite No. ¹	Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine, journal, serial, symposium, catalog, etc.), date, page(s), volume-issue number(s), publisher, city and/or country where published.	T ⁶
	FA	OIVANEN, M., <i>et al.</i> , "Additional evidence for the exceptional mechanism of the acid-catalysed hydrolysis of 4-oxypyrimidine nucleosides: hydrolysis of 1-(1-alkoxyalkyl)uracils. Seconucleosides. 3'-C-alkyl nucleosides and nucleosides 3',5'-cyclic monophosphates," <i>J. Chem. Soc. Perkin Trans.</i> , 1994, 2, 309-314.	
	FB	ONG, S.P., <i>et al.</i> , "Synthesis of 3'-C-methyl adenosine and 3'-C-methyluridine diphosphates and their interaction with the ribonucleoside diphosphate reductase from <i>Corynebacterium nephridii</i> ," <i>Biochemistry</i> , 1992, 31, 11210-11215.	
	FC	PAN-ZHOU X-R., <i>et al.</i> , "Differential effects of antiretroviral nucleoside analogs on mitochondrial function in HepG2 cells," <i>Antimicrob Agents Chemother</i> 2000, 44(no.3), 496-503.	
	FD	ROSENTHAL, A., <i>et al.</i> , "Branched-chain sugar nucleosides. Synthesis of 3'-C-ethyl (and 3'-C-butyl)uridine," <i>Carbohydrate Research</i> , 1980, 79, 235-242.	
	FE	SAMANO, V., <i>et al.</i> , "Nucleic acid related compounds. 77. 2',3'-didehydro-2',3'-dideoxy-2'-(and 3')-methyl nucleosides via [3,3]-sigmatropic rearrangements of 2'-(and 3')-methylene-3'-(and 2')-O-thiocarbonyl derivatives and radical reduction of a 2'-chloro-3'-methylene analogue," <i>Can. J. Chem.</i> , 1993, 71, 186-191.	
	FF	SAMANO, V., <i>et al.</i> , "Synthesis and radical-induced ring-opening reactions of 2'-deoxyadenosine-2'-spirocyclopropane and its uridine analogs. Mechanistic probes for ribonucleotide reductases," <i>J. Am. Chem. Soc.</i> , 1992, 114, 4007-4008.	
	FG	SCHMIT, C. <i>et al.</i> , "The effects of 2'- and 3'-alkyl substituents on oligonucleotide hybridization and stability," <i>Biorganic & Medicinal Chemistry Letters</i> , 1994, 4(No. 16), 1969-1974.	
	FH	SERAFINOWSKI, P.J., <i>et al.</i> , "New method for the preparation of some 2'- and 3'-trifluoromethyl-2'-3'-dideoxyuridine derivatives," <i>Tetrahedron</i> , 1999, 56(No. 2), 333-339.	
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	FJ	SOMMADOSSI J-P., <i>et al.</i> , "Comparison of cytotoxicity of the (-) and (+)-enantiomer of 2',3'-dideoxy-3'-thiaeytidine in normal human bone marrow progenitor cells," <i>Biochemical Pharmacology</i> , 1992, 44:1921-1925.	
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	FL	TRITSCH, D., <i>et al.</i> , "3'-β-ethynyl and 2'-deoxy-3'-β-ethynyl adenosines: first 3'-β-branched-adenosines substrates of adenosine deaminase," <i>Biorganic & Medicinal Chemistry Letters</i> , 2000, 10, 139-141.	

Examiner Signature

Howard V. Owens, Jr.

Date Considered

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Sheet	7	of	7
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Complete if Known

Application Number	09/834,596
Filing Date	April 13, 2001
First Named Inventor	Watanabe <i>et al.</i>
Group Art Unit	1623
Examiner Name	Howard V. Owens, Jr.
Attorney Docket Number	08841.105037 PHAR 2020

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Sheet 1 of 3

Application Number	09/834,596
Filing Date	13 April 2001
First Named Inventor	Watanabe, K.
Group Art Unit	1614
Examiner Name	Not Assigned
Attorney Docket Number	PHARM1

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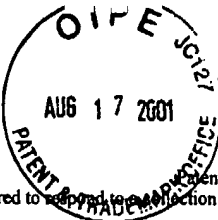
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Substitute for form 1449B/PTO		Application Number	09/834,596
INFORMATION DISCLOSURE STATEMENT BY APPLICANT (use as many sheets as necessary)		Filing Date	04/13/01
		First Named Inventor	WATANABE
		Group Art Unit	1614
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Ho		Fedorov, I., et al., XIII International Round Table: Nucleosides, Nucleotides and Their Biological Applications. Montpellier, France, September 6-10, 1998. Poster 35.	
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Substitute for form 1449B/PTO		Application Number	09/834,596
INFORMATION DISCLOSURE		Filing Date	04/13/01
STATEMENT BY APPLICANT		First Named Inventor	WATANABE
(use as many sheets as necessary)		Group Art Unit	1614
		Examiner Name	Not Assigned
		Attorney Docket Number	PHARM1
Sheet	3	of	3

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AO		Pudlo, J., Nucleosides & Nucleotides, 11(2-4), 279-93 (1992)	
		Fiandor, J., et al., Nucleosides & Nucleotides, 8(5&6), 1107-8 (1989)	
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WJO		Qiu, Y., et al., J. Med. Chem., 1998, 41:10-23	
AO		Sekiyama, T., J. Med. Chem., 1998, 41:1284-98	

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